Refine Search

Search Results -

Terms	Documents
pyrogallol adj8 cyclic and L6	5

Database:

US Pre-Grant Publication Full-Text Database
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JPO Abstracts Database
Derwent World Patents Index

IBM Technical Disclosure Bulletins

Search:

L7		Refine Search
J	Recall Text Clear	Interrupt

Search History

DATE: Thursday, January 11, 2007 Purge Queries Printable Copy Create Case

Set Name side by side	Query	Hit Count	Set Name result set
DB = PGPB, US	SPT,USOC,EPAB,JPAB,DWPI,TDBD; PLUR=	YES; OP=ADJ	•
<u>L7</u>	pyrogallol adj8 cyclic and 16	5	<u>L7</u>
<u>L6</u>	L5 and (treat\$7 or infect\$7)	76	<u>L6</u>
<u>L5</u>	L3 and (hiv or aids)	80	<u>L5</u>
<u>L4</u>	L3 and phenyl tetramer	1	<u>L4</u>
<u>L3</u>	L2 and cyclic	192	<u>L3</u>
<u>L2</u>	L1 and (514/\$ or 562/\$)	469	<u>L2</u>
<u>L1</u>	pyrogallol	13290	L1

END OF SEARCH HISTORY

Hit List

First Hit Clear Generate Collection Print Fwd Refs Bkwd Refs Generate OACS

Search Results - Record(s) 1 through 5 of 5 returned.

☐ 1. Document ID: US 20050288228 A1

L7: Entry 1 of 5

File: PGPB

Dec 29, 2005

PGPUB-DOCUMENT-NUMBER: 20050288228

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20050288228 A1

TITLE: Bile-acid conjugates for providing sustained systemic concentrations of

drugs

PUBLICATION-DATE: December 29, 2005

INVENTOR - INFORMATION:

NAME

CITY .

STATE

COUNTRY

Cundy, Kenneth C.

Redwood City

CA

US

Gallop, Mark A.

Los Altos

CA

US

US-CL-CURRENT: <u>514/12</u>; <u>514/169</u>, <u>514/44</u>

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw, De

☐ 2. Document ID: US 20020142998 A1

L7: Entry 2 of 5

File: PGPB

Oct 3, 2002

PGPUB-DOCUMENT-NUMBER: 20020142998

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020142998 A1

TITLE: Bile-acid conjugates for providing sustained systemic concentrations of

drugs

PUBLICATION-DATE: October 3, 2002

INVENTOR - INFORMATION:

NAME

CITY

STATE

COUNTRY

Cundy, Kenneth C.

Gallop, Mark A.

Redwood City

Los Altos

CA CA US US

US-CL-CURRENT: <u>514/169</u>; <u>514/182</u>, <u>552/515</u>, <u>552/540</u>

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw. De

□ 3. Document ID: US 20020099041 A1

L7: Entry 3 of 5

File: PGPB

Jul 25, 2002

PGPUB-DOCUMENT-NUMBER: 20020099041

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020099041 A1

TITLE: Bile-acid derived compounds for enhancing oral absorption and systemic

bioavailability of drugs

PUBLICATION-DATE: July 25, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

Gallop, Mark A.

Los Altos

CA

US

Cundy, Kenneth C.

Redwood City

CA

US

US-CL-CURRENT: <u>514/169</u>

.	Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw, De
									•				

☐ 4. Document ID: US 7144877 B2

L7: Entry 4 of 5

File: USPT

Dec 5, 2006

US-PAT-NO: 7144877

DOCUMENT-IDENTIFIER: US 7144877 B2

TITLE: Bile-acid derived compounds for enhancing oral absorption and systemic

bioavailability of drugs

PRIOR-PUBLICATION:

DOC-ID

DATE

US 20020099041 A1

July 25, 2002

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sanyances	Wildhinank	Claims	KWIC	Draw, De
						-,						

☐ 5. Document ID: US 6984634 B2

L7: Entry 5 of 5

File: USPT

Jan 10, 2006

US-PAT-NO: 6984634

DOCUMENT-IDENTIFIER: US 6984634 B2

TITLE: Bile-acid conjugates for providing sustained systemic concentrations of

drugs

PRIOR-PUBLICATION:

DOC-ID

US 20020142998 A1

DATE

October 3, 2002

Full	Title Citation	Front	Review	Classification	Date	Reference	SECT	encea	Alicial	Haries	Claims	KWIC	Draw, De
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	pyrogall	ol a	dj8 c	yclic a	nd L	6						5	

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Previous Page Next Page Go to Doc#

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chain nodes:

14 15 16 17 18 32 33 34 35 36 50 51 52 53 54 68 69 70 71 72 ring nodes:

1 2 3 4 5 6 7 8 9 10 11 12 13 19 20 21 22 23 24 25 26 27 28 29 30 31 37 38 39 40 41 42 43 44 45 46 47 48 49 55 56 57 58 59 60 61 62 63 64 65 66 67 chain bonds :

3-16 4-15 5-17 7-18 10-13 13-14 21-34 22-33 23-35 25-36 28-31 31-32 39-52 40-51 41-53 43-54 46-49 49-50 57-70 58-69 59-71 61-72 64-67 67-68 ring bonds :

1-2 1-6 2-3 2-31 3-4 4-5 5-6 6-13 7-8 7-12 8-9 9-10 10-11 11-12 13-38 19-20 19-24 20-21 20-67 21-22 22-23 23-24 24-31 25-26 25-30 26-27 27-28 28-29 29-30 37-38 37-42 38-39 39-40 40-41 41-42 42-49 43-44 43-48 44-45 45-46 46-47 47-48 49-56 55-56 55-60 56-57 57-58 58-59 59-60 60-67 61-62 61-66 62-63 63-64 64-65 65-66 exact/norm bonds :

2-31 3-16 4-15 5-17 6-13 13-38 20-67 21-34 22-33 23-35 24-31 39-52 40-51 41-53 42-49 49-56 57-70 58-69 59-71 60-67

exact bonds:

7-18 10-13 13-14 25-36 28-31 31-32 43-54 46-49 49-50 61-72 64-67 67-68 normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-12 8-9 9-10 10-11 11-12 19-20 19-24 20-21 21-22 22-23 23-24 25-26 25-30 26-27 27-28 28-29 29-30 37-38 37-42 38-39 39-40 40-41 41-42 43-44 43-48 44-45 45-46 46-47 47-48 55-56 55-60 56-57 57-58 58-59 59-60 61-62 61-66 62-63 63-64 64-65 65-66

Match level:

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom 12:Atom 13:Atom 14:CLAS\$15:CLAS\$16:CLAS\$17:CLAS\$18:CLAS\$19:Atom 20:Atom 21:Atom 22:Atom 23:Atom 24:Atom 25:Atom 26:Atom 27:Atom 28:Atom 29:Atom 30:Atom 31:Atom 32:CLAS\$33:CLAS\$34:CLAS\$ 35:CLAS\$36:CLAS\$37:Atom 38:Atom 39:Atom 40:Atom 41:Atom 42:Atom 43:Atom 44:Atom 45:Atom 46:Atom 47:Atom 48:Atom 49:Atom 50:CLAS\$51:CLAS\$52:CLAS\$53:CLAS\$54:CLAS\$55:Atom 56:Atom 57:Atom 58:Atom 59:Atom 60:Atom 61:Atom 62:Atom 63:Atom 64:Atom 65:Atom 66:Atom 67:Atom 68:CLAS\$69:CLAS\$70:CLAS\$71:CLAS\$72:CLAS\$

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SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FILE 'CAPLUS' ENTERED AT 12:12:07 ON 11 JAN 2007

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L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1

STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Structure attributes must be viewed using STN Express query preparation.

=> s 11 full

REG1stRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress... Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

23 ANSWERS

FULL SEARCH INITIATED 12:12:39 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 161 TO ITERATE

100.0% PROCESSED 161 ITERATIONS

SEARCH TIME: 00.00.01

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L3 5 L2

=> s 13 and py<2003 22868731 PY<2003

L4 2 L3 AND PY<2003

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2002:428850 CAPLUS

DOCUMENT NUMBER: 137:6006

TITLE: Preparation of Calixarenes as Anti-viral compounds

INVENTOR(S):
Harris, Stephen J.

PATENT ASSIGNEE(S): Aids Care Pharma Limited, Ire.

SOURCE: PCT Int. Appl., 44 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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PATENT NO.
                                                       APPLICATION NO.
                               KIND
                                        DATE
                                                                                    DATE
                                        20020606
                                                      WO 2001-IE150
      WO 2002044121
                                A1
                                                                                    20011130 <--
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                CN, CO, CR, CU, CZ, CZ, DE, DE, DK, DK, DM, DZ, EE, EE, ES, FI, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK, SL, TJ, TM,
                TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ,
                MD, RU, TJ, TM
           RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,
                CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,
                BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                                        20020611
                                                      AU 2002-20992
      AU 2002020992
                                A5
                                                                                    20011130 <--
      EP 1345884
                                A1
                                        20030924
                                                      EP 2001-998526
                                                                                    20011130
               AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
                IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
PRIORITY APPLN. INFO.:
                                                       IE 2000-983
                                                                                A 20001201
                                                       WO 2001-IE150
                                                                                W 20011130
OTHER SOURCE(S):
                               CASREACT 137:6006; MARPAT 137:6006
GΙ
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* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

- AB Title compds. I [R1 = OCH2CO2K, OCH2CO2H or OCH2CONH2; R2 = R1 or NO2; R3 = H, 2-HO2CCH2OC6H4, or 4-XC6H4 where X = halo (preferably F or Br); R4 = H or halo (preferably Br)] are prepared and disclosed as antiviral agents. Thus, II was prepared in four steps via cyclocondensation 4-fluorobenzaldehyde with pyrogallol and subsequent bromination, O-alkylation with Et bromoacetate and hydrolysis with KOH. II possessed a therapeutic index (TC50/EC50 µm) of 4,000. I were found to have an additive effect when administered with AZT, and therefore, the compds. are useful as pharmaceutical compns. in the treatment of AIDS.
- IT 433334-86-2P 433334-87-3P 433334-88-4P

433334-94-2P 433334-95-3P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(intermediates; preparation and antiviral activity of calixarenes as anti-AIDS agents)

RN 433334-86-2 CAPLUS

CN Pentacyclo[19.3.1.13,7.19,13.115,19]octacosa-1(25),3,5,7(28),9,11,13(27),1 5,17,19(26),21,23-dodecaene-4,5,6,10,11,12,16,17,18,22,23,24-dodecol, 2,8,14,20-tetrakis(4-fluorophenyl)- (9CI) (CA INDEX NAME)

RN 43334-87-3 CAPLUS
CN Pentacyclo[19.3.1.13,7.19,13.115,19]octacosa-1(25),3,5,7(28),9,11,13(27),1
5,17,19(26),21,23-dodecaene-4,5,6,10,11,12,16,17,18,22,23,24-dodecol,
25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4-fluorophenyl)- (9CI) (CA
INDEX NAME)

OEt

RN 433334-94-2 CAPLUS

CN Pentacyclo[19.3.1.13,7.19,13.115,19]octacosa-1(25),3,5,7(28),9,11,13(27),1 5,17,19(26),21,23-dodecaene-4,5,6,10,11,12,16,17,18,22,23,24-dodecol, 2,8,14,20-tetrakis(4-bromophenyl)- (9CI) (CA INDEX NAME)

RN 433334-95-3 CAPLUS

CN Pentacyclo[19.3.1.13,7.19,13.115,19]octacosa-1(25),3,5,7(28),9,11,13(27),1 5,17,19(26),21,23-dodecaene-4,5,6,10,11,12,16,17,18,22,23,24-dodecol, 25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4-bromophenyl)- (9CI) (CA INDEX NAME)

PAGE 1-A

●12 K

PAGE 1-A

$$\begin{array}{c|c} & & & & \\ & &$$

PAGE 2-B

^{_}NH₂

// \ NH2 PAGE 3-A

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

1995:994163 CAPLUS

DOCUMENT NUMBER:

124:55584

TITLE:

Preparation of calixarene-based compounds having antibacterial, antifungal, anticancer, and anti-HIV

activity

INVENTOR(S):

Harris, Stephen J.

PATENT ASSIGNEE(S):

Ire.

SOURCE:

PCT Int. Appl., 148 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent English

LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

. PATENT NO.				KIN	KIND DATE				APPLICATION NO.					DATE			
WO 9519974 WO 9519974					A2 19950727				WO 1995-IE8					19950124 <			
WO			n i i	DD	A3		1995			CN	C7	DE	DIZ	ът	CP	1111	TD.
					RO,		BY, US	CA,	Cn,	CN,	CZ,	υĿ,	DK,	ΓI,	GD,	no,	UP,
	RW:	ΑT,	BE,	•	•	•	FR,	GB,	GR,	IE,	LU,	NL,	SE,	GA,	ML,	NE,	SN,
		TD,	TG														
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PRIORIT	Y APP	LN.	INFO	.:						IE 1	994-	57			A 1	9940	124
									1	WO 1	995-:	IE8			A 1	9950	124

OTHER SOURCE(S):

MARPAT 124:55584

For diagram(s), see printed CA Issue. GI

- Calixarene-based compds., which are calixarenes or oxacalixarenes, acyclic AB phenyl-formaldehyde oligomers, cyclotriveratrylene derivs., cyclic tetrameric resorcinol-aldehyde derivs. known as Hogberg compds. and cyclic tetrameric pyrogallol-aldehyde derivs., are prepared For example, calixarenes or oxacalixarenes are represented by general formula [I; n + m = 3-8; m = 0-3; n = 0-8; R1 = H, halo, hydrocarbyl, aryl, (un)substituted hydrocarbylaryl, NO2, SO3M1; wherein M1 = alkali metal, SO3H; R1 = OR2; wherein R2 = CH2CO2R3, CH2CO2Mp/p, CH2CONR4R5; wherein R3 = (un) substituted alkyl; M = metal, ammonium ion; p = the charge on themetal ion; R4 or R5 may be the same or different, or both may be part of amino acid ester of poly(amino acid ester) or one or more of the same or different amino acids or part of a cyclic polyene antibiotic/antifungal drug or part of a cyclic nitrogen heterocycle; X = halo, NO2, CO2H, cyano, other electron withdrawing group]. Thus, n-butyraldehyde and pyrogallol in a 1:4 mixture of 37% aqueous HCl and EtOH was refluxed under N for 90 min to give a cyclic tetramer (II; R = X = H), which was brominated with Br in CHCl3 to II (R = H, X = Br) and etherified with Et bromoacetate in the presence of K2CO3 in refluxing acetone to give II (R = CH2CO2Et, X = Br). The latter compound was saponified with KOH in refluxing EtOH , acidified with aqueous HCl, and treated with 25% aqueous NH4OH to give II (R = CH2CO2-NH4+, X
 - Br). The latter compound in vitro inhibited the infection of C8166 cells with HIV-2, SIV (Simian immunodeficiency virus), and HIV-1 with EC50 of 10, 20, and 0.03 μ M.
- ΙT 171799-80-7P 171799-81-8P 171799-82-9P 171799-89-6P 171799-90-9P 171799-91-0P 171799-95-4P 171799-96-5P 171799-97-6P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of calixarene-based compds. having antibacterial, antifungal, anticancer, and anti-HIV activity)

RN

CN

''',2''''',2''''',2'''''''-[[25,26,27,28-tetrabromo-2,8,14,20-(10-chloro-9-anthracenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacosa-1(25),3,5,7(28),8,11,13(27),15,17,19(26),21,23-dodecaene-4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis-, dodecapotassium salt (9CI) (CA INDEX NAME)

PAGE 4-A

●12 K

PAGE 4-A

PAGE 4-A

●12 NH3

●12 K

●12 NH3

●12 K

●12 NH3